

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
14 July 2005 (14.07.2005)

PCT

(10) International Publication Number
WO 2005/063690 A1

(51) International Patent Classification⁷: **C07C 235/40**,
C07D 271/06, C07C 235/12, 235/34, 235/26, 235/06,
23/08, C07D 213/61, A61K 31/235, 31/4245, 31/165,
31/44, A61P 29/00

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(21) International Application Number:
PCT/US2004/042691

(22) International Filing Date:
17 December 2004 (17.12.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/531,643 22 December 2003 (22.12.2003) US
60/539,637 28 January 2004 (28.01.2004) US
60/624,958 4 November 2004 (04.11.2004) US

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(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,
AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
ZW.

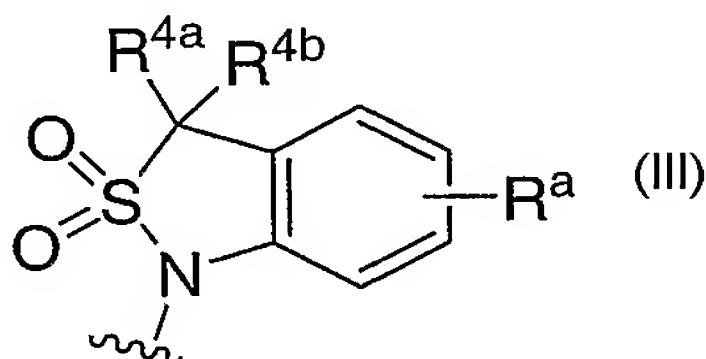
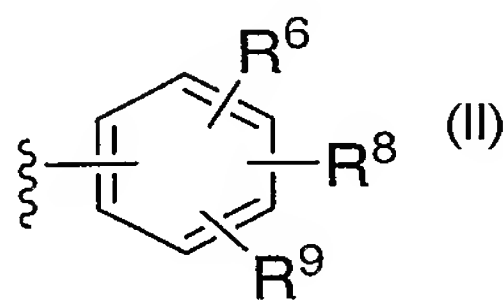
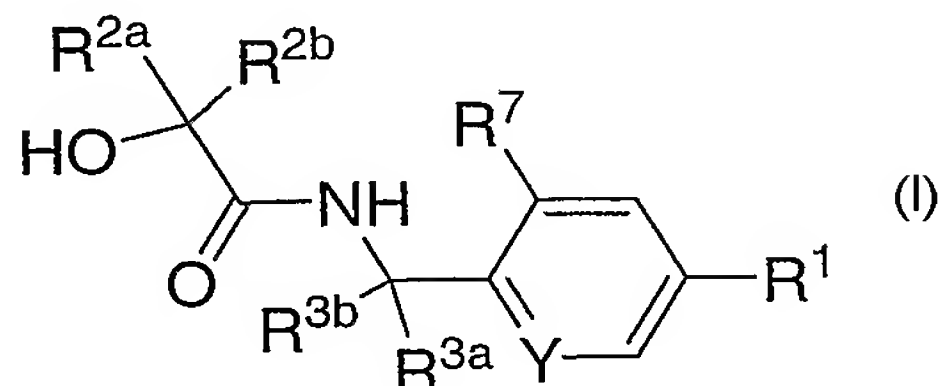
(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,
ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI,
FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO,
SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN,
GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: ALPHA-HYDROXY AMIDES AS BRADYKININ ANTAGONISTS OR INVERSE AGONISTS



(57) Abstract: α -Hydroxy amide derivatives of the general formula (I) are bradykinin B1 antagonists or inverse agonists useful in the treatment or prevention of symptoms such as pain and inflammation associated with the bradykinin B1 pathway. R^{2a} is selected from (1) a group selected from R^a . (2) $(CH_2)_nNR^bC(O)R^a$. (3) $(CH_2)_nNR^bSO_2R^d$. (4) $(CH_2)_nNR^bCO_2R^a$. (5) $(CH_2)_k$ -heterocycle optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, OR^a , SR^a , C_{1-4} alkyl and C_{1-3} haloalkyl wherein said heterocycle is (a) a 5-membered heteroaromatic ring having a ring heteroatom selected from N, O and S, and optionally having up to 3 additional ring nitrogen atoms wherein said ring is optionally benzo-fused; or (b) a 6-membered heteromatic ring containing from 1 to 3 ring nitrogen atoms and

N-oxydes thereof. Wherein said ring is optionally benzo-fused. (6) $(CH_2)_kCO_2R^a$. and (7) $(CH_2)_kC(O)NR^bR^c$. R^{2b} is OH or a group selected from R^{2a} ; or R^{2a} and R^{2b} together with the carbon atom to which they are attached form a 3- to 7-membered carbocyclic ring optionally substituted with 1 to 4 groups independently selected from halogen, OR^a , C_{1-4} alkyl and C_{1-4} haloalkyl;